Rare Complications of Silica Dust Exposure

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INTRODUCTION

Silicosis is a diffuse irreversible interstitial lung disease caused by continuous inhalation of crystalline silica (SiO₂) leading to lung fibrosis (1). Because of the absence of a definitive silicosis treatment, prevention of the disease is the main concern. To prevent the disease, regular close monitoring of the workplaces and routine screening of the workers are required, in addition to the establishment of effective workplace regulations (2). In this paper, we report two cases of rare complications of silica dust exposure.

CASE REPORTS

Case 1

A 34-year-old male, who used to work in glass grinding, complained of exertional dyspnea and dry cough for 1 year. Three months before, dyspnea worsened to occur with mild exertion, and the patient developed cyanosis. On examination, he was cyanotic with oxygen saturation 84% on room air. A high-resolution computed tomography (HRCT) chest scan showed diffuse ground-glass opacification, septal thickening, and areas of a “crazy-paving” pattern (Fig. 1a). Bronchoalveolar lavage (BAL) from the right lower lobe was milky white (Fig. 1b), and transbronchial lung biopsies (TBLB) form the right lower lobe revealed intra-alveolar amorphous proteinaceous material associated with hyperplastic alveolar macrophages and interstitial inflammation (Fig. 1c). The intra-alveolar material was periodic acid Schiff (PAS) stain-positive (Fig. 1d), confirming the diagnosis of alveolar proteinosis.

The whole hole-lung lavage was performed for both lungs sequentially.

The more severely affected lung, as detected by a CT scan, was lavaged first, and the other after 24–48 hr. Accordingly, the patient was intubated with a double-lumen endotracheal tube, and after 15 min of ventilation with 100% O₂, one lung was lavaged with sterile isotonic saline at 37°C. The volume used for each filling was 1000 ml, and then the lung is left to drain by gravity. The filling and drainage was repeated till the effluent is clear, and 10 L of saline was required for each lung.

Thereafter, the patient’s symptoms and oxygen saturation had improved, and his oxygen saturation was 93%.

Case 2

A 31-year-old male, who used to work in sandblasting, ex-smoker, complained of exertional dyspnea for 1 year. One month before, dyspnea worsened to occur with mild exertion, and the patient developed night fever and night sweats with loss of weight and appetite, and productive yellowish sputum. On examination, he was cachectic and run a fever 38°C, with bilateral infraclavicular crepitations. A chest CT scan showed diffuse ground-glass opacification, septal thickening, and areas of a “crazy-paving” pattern (Fig. 2a). Bronchoalveolar lavage (BAL) from the right lower lobe was milky white (Fig. 1b), and transbronchial lung biopsies (TBLB) form the right lower lobe revealed intra-alveolar amorphous proteinaceous material associated with hyperplastic alveolar macrophages and interstitial inflammation (Fig. 1c). The intra-alveolar material was periodic acid Schiff (PAS) stain-positive (Fig. 1d), confirming the diagnosis of alveolar proteinosis. The whole hole-lung lavage was performed for both lungs sequentially.

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namide, and ethambutol was initiated, and the patient continued therapy for 6 months.

**DISCUSSION**

Silicosis is an incurable occupational interstitial lung disease caused by free crystalline silica inhalation and its accumulation in the lung interstitium. Variable forms of the disease can be identified, but mainly three can be identified: chronic/classic, accelerated, and acute. Alveolar silicoproteinosis is one of the acute complications of silica dust exposure with subsequent alveolar filling with proteinaceous material (3). Massive inhalation of silica dust leads to shortness of breath, productive cough, chest pain and loss of weight, fever, and fatigue (4).

The standard treatment for alveolar silicoproteinosis is whole lung lavage, which removes large amounts of silica dust and inflammatory cells. Thereafter, it relieves pulmonary symptoms and improves oxygenation (5).

Exposure to silica, even without initiation of silicosis disease is associated with a high-risk predisposition of TB, which has been reported as 1.9% per year. A reduction in silica dust exposure, continuous medical surveillance, and TB screening in high-risk occupations and securing anti-TB treatment are strategies to minimize the rate of silicotuberculosis among workers or employees (6).

**CONCLUSION**

In the current cases, we have confirmed a rare diagnosis of acute silicoproteinosis and silicotuberculosis. Given the patients’ occupational history of glass grinding and sandblasting and known high-risk exposure to respirable silica dust in their professions, we have considered a possible occupational etiology. Finally, this was confirmed by clinical, cytological, and histopathological findings.

**Informed Consent:** Written informed consent was obtained from patients who participated in this study.

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**REFERENCES**